GUIDELINE FOR THE EVALUATION OF MOBILE C-ARM FLUOROSCOPIC X-RAY EQUIPMENT



State of Utah
Department of Environmental Quality
Division of Radiation Control

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DRC Inspection Program Objective

The overall objective of the Division of Radiation Control (DRC) x-ray inspection program is **6** reduce the likelihood that individuals will be exposed to unnecessary radiation. In the case **6** registrants using x-ray equipment in the healing arts, patient exposure is of concern and proper equipment performance is essential. Registrants are required to demonstrate that the equipment satisfies the appropriate regulatory standards for calibration and performance.

Purpose of Guideline

The intent and purpose of this document is to provide users of **mobile c-arm fluoroscopic x-ray equipment** guidelines for the documentation required to demonstrate to the DRC that the x-ray equipment satisfies the regulatory standards under clinical use conditions.

X-ray Equipment Performance and Calibration

The registrant is to document that the following requirements are met.

- 1) Adequate total filtration is present.
- 2) The fluoroscopy timer terminates the exposure or produces an audible signal at the end of a five minute accumulative time interval.
- 3) During fluoroscopy, the x-ray field collimation and alignment with the image intensifier (II) is appropriate.
- 4) Fluoroscopic exposure rates do not exceed the regulatory standards.
- Patient exposure information has been obtained for simulated clinical conditions and is posted where it is readily available to the physician during the fluoroscopic procedure.

The following examples are presented as guidance for what will be considered an adequate evaluation, with support documentation, to demonstrate compliance:

1) Adequate Filtration

Demonstration of adequate filtration shall be accomplished by showing that the half value layer (HVL) exceeds the minimum regulatory standard. For example, at a measured kVp value of 80, the HVL is to be equal to or greater than 2.3 mm aluminum. This can be demonstrated by:

a) Measuring the in air exposure when different thicknesses of aluminum intercept the x-ray beam and determining the HVL value; or

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b) Measuring the exposure at 80 kVp with and without a 2.5 mm aluminum absorber intercepting the x-ray beam and showing that the ratio of the two exposure values exceeds 0.5.

(Documentation shall include a listing of measured exposure values and associated thicknesses of aluminum.)

2) Fluoroscopic Timer

The fluoroscopy timer terminates the exposure or produces an audible signal at the end of a five minute accumulative time interval.

(Documentation shall indicate that the timer was evaluated and that a conclusion was reached whether the exposure was terminated at the end of five minutes or an audible signal occurred.)

3) Fluoroscopic X-ray Field Collimation and Alignment

For c-arm mobile fluoroscopic x-ray equipment, the x-ray field is to be aligned with and collimated to the effective field size of the II tube within the regulatory standards. Annual tests are to be carried out to insure that the above condition is met.

(Documentation is available that indicates tests were performed demonstrating that the maximum field size of the x-ray field is confined to the II within the regulatory standards for the largest effective II size that is available. Evidence of collimation observed on the monitor will be deemed adequate.)

4) <u>Limits on Exposure Rates</u>

Exposure rates are to be evaluated to insure that the regulatory limits are not exceeded Documentation will indicate whether the fluoroscopic unit has a manual mode, automatic brightness control (ABC) mode, high level control (HLC) mode, the sizes of II available, and if the unit was manufactured on or before 05/19/95. (NOTE: Different limits apply for units manufactured before and after this date.)

Example 1: A mobile c-arm fluoroscopic unit is used in a hospital surgical facility. The unit has three different ABC modes of operation and a nine inch II effective field size. For all three ABC modes, both the kVp and mA are automatically chosen. For a given patient, the specific kVp and mA values and the corresponding exposure rate depend on which ABC mode is used. With a lead absorber intercepting the x-ray beam, the exposure rate is measured, at 30 cm from the II, in the fluoroscopic mode which will produce the maximum exposure rate.

(Documentation will indicate the ABC mode used to measure the maximum exposure rate, the position relative to the II where the measurements were made, the value obtained, and the

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corresponding kVp and mA values. A statement to the effect that the unit satisfies the regulatory limits, and that the evaluation was adequate to insure that the mode usedproduced the maximum rate, is to be included.)

5) Patient Exposure Information

The Utah Radiation Control Rules require patient exposure information be readily available to the attending physician during a fluoroscopic procedure. The policy of the Division is that such information is to be available for mobile c-arm units used in the irradiation of either the pelvis, abdominal, or upper GI regions. Information shall be in a format such that entrance skin exposure rates can be readily determined from a knowledge of the fluoroscopic kVp and mA values. The clinical situation for which the exposure information is applicable and the geometrical location of the entrance skin relative to the x-ray source is to be indicated.

Example formats deemed acceptable to the Division are included. (Note: The Division recognizes that other formats are possible which would meet the intent of the regulations.)

Example 1: A mobile unit is used in surgery. It has "normal", "normal 1/2 dose," "pulsed normal," and "snap shot" ABC modes. The snap shot mode is used for surgical procedures involving extremities. For retrograde cystogram procedures, the normal mode is used in an under table configuration.

X-ray Unit: Date:	
Date:	
Entrance Skin Exposure Rate (mR/min)	
at the table top	
mA	A
	_ in
kVp table	V
tube	

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Example 2: The mobile c-arm unit is used in surgery for laparoscopic cholecystectomy (removal of the gallbladder) procedures. The unit has several ABC modes, but is only used in the "normal" mode. It is used in both under table and lateral geometrical configurations.

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	t:						
		Entrance Skin Exposure Rate (mR/min) at the table top mA				Image Intensifier	
						1 _	_ in.
kVp						table	
						tube	
		•	•			1	
— X-ray Uni	cut & post						
			Skin Expo @ pt. A mA		(R/min)	Pt.A tube	Image
						x	Intensifier
	70					table	
kVp	80						
	90					pt. A is 30 cm. fro image intensifier	om
	100						

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